FDA 510(k) Premarket Notification

K024170 L1 SYSTEM

CLEARED

Apr 14, 2003
117 days to decision

K024170 · Product code: **FMK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k024170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Dec 18, 2002
Decision date	Apr 14, 2003
Days to decision	117 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pelikan Technologies, Inc.**
Location	Palo Alto, CA, US
Contact	JACK GREEN
510(k) history	2 submissions · 2 cleared · 2003-2004